INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
(Not for submission under 37 CFR 1.99)

Application Number		09444774
Filing Date		2000-11-22
First Named Inventor	Mikur	rak
Art Unit		3622
Examiner Name	A. Du	iran
Attorney Docket Number		60021-334801

					U.S.	PATENTS			Remove	
Examiner Initial*	Cite No	Patent Number	Kind Code ¹	1 Issue Date		Name of Pate of cited Docu	entee or Applicant ment	Releva	Columns,Lines when int Passages or Rele s Appear	
	1	5974395		1999-1	0-26	Belini et al.				
If you wisl	h to a	l dd additional U.S. Pate	nt citatio	n inform	ation pl	ease click the	Add button.	_	Add	
			U.S.P	ATENT	APPLI	CATION PUB	LICATIONS		Remove	
Examiner Initial*	Cite No	Publication Number	Kind Code ¹	Publica Date	ition	Name of Patentee or Applicant of cited Document		Releva	Columns,Lines when int Passages or Rele s Appear	
	1									
If you wisl	h to a	dd additional U.S. Publi						d button		
				FOREIG	3N PAT	ENT DOCUM	ENTS		Remove	
Examiner Initial*	Cite No	Foreign Document Number ³	Country Code ²		Kind Code ⁴	Publication Date	Name of Patente Applicant of cited Document	e or	Pages,Columns,Line: where Relevant Passages or Relevan Figures Appear	
	1									
If you wisl	h to a	l dd additional Foreign P	atent Do	cument	citation	information pl	lease click the Add	button	Add	
			NON	I-PATE	NT LITE	RATURE DO	CUMENTS		Remove	
Examiner Cite Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc), date, pages(s), volume-issue number(s), publisher, city and/or country where published.					Ţŧ					

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99) Art Unit Examiner

Application Number		09444774
Filing Date		2000-11-22
First Named Inventor Mikur		ak
Art Unit		3622
Examiner Name A. Du		iran
Attorney Docket Number		60021-334801

	1	SODHI, M., "Cyberspace: Internet-Enabled Collaborative Planning," OR/MS Today, June 1998, (1 page)						
	2	SAP® Advanced Planner and Optimizer -Collaborative Planning, October 1999, pp. 1-20, SAP AG, Walldorf Germany						
If you wish to add additional non-patent literature document citation information please click the Add button Add								

If you wish to add additional non-patent literature document citation information please click the Add button Add

EXAMINER SIGNATURE

Examiner Signature Date Considered

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through a citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

See Kinz Codes of USPTO Potent Documents at least USETO_COLV or MEDE 901.04. * Enter of this that issued the document, by the load-lester code (NEO) Searchest 31.0.* * Enter of the paperes period for countent, the ordinated or the parent be ready or the Emperor many process the serial number of the patent document.
*Viole of coursent by the appropriate symbols as indicated on the document under WPO Standard 31.16 if passible. * Applicant is to place a check mark here if English tanguage seriations as although the page of the page of the page of the passible. * Applicant is to place a check mark here if

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

	Application Number		09444774		
	Filing Date		2000-11-22		
	First Named Inventor	Mikur	ak		
Art Unit			3622		
	Examiner Name	A. Du	Duran		
	Attorney Docket Number		60021-334801		

CERTIFICATION STATEMENT

Please see 37 CFR 1.97 and 1.98 to make the appropriate selection(s):

That each item of information contained in the information disclosure statement was first cited in any communication of from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. See 37 CFR 197(e)(1).

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquiry, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 1/56(c) more than three months prior to the filing of the information disclosure statement, See 37 CFR 1/97(c).

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.

Dena VanDeVoort Ehrich

□ None

Name/Print

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

Registration Number

form or the signature.			
Signature	/DenaVanDeVoortEhrich/	Date (YYYY-MM-DD)	2006-06-29

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for life and by the USPTO to process) an application. Confidentially is governed by \$5 U.S. C. 12 and 37 CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, u.S. Operament of Commence, P. 0. Box 1450, Alexandris, V.2.231-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.2.2313-1450.

Privacy Act Statement

The Privacy Act of 1974 (P. L. 95.79) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that. (1) the general authority for the collection of this information is \$5 U.S. C. 2(b)(2); (2) furnishing of the information solicide that.) (1) the general authority for the collection of this information is used by the U.S. Patient and Trademan Koffice is to process and/or examine your submission related to a patient application or patient. If you do not furnish the requested in the contraction of the contraction of the patient o

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiations.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record pertains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, pursuant to 5 U.S. C. 552(m).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
- 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designe, during an inspection of records conducted by GSA a part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S. C. 294 and 2906. Such Adeckouse shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the againston pursuant to 35 U.S. C. 12(b) or issuance of a patient pursuant to 35 U.S. C. 151. Further, a record may be disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record vals filed in an application which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application pare to public insepections or an issued patient.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.